

1) Publication number:

0 423 705 A3

·(12)

## **EUROPEAN PATENT APPLICATION**

2) Application number: 90119803.6

2 Date of filing: 16.10.90

(9) Int. Cl.<sup>5</sup>: **C07B 55/00**, C07C 323/32, C07C 317/32, C07C 323/39, C07C 215/34, C07C 213/10, C07D 263/06

Priority: 20.10.89 IT 2207589

Date of publication of application:24.04.91 Bulletin 91/17

Designated Contracting States:
 AT BE CH DE DK ES FR GB GR IT LI LU NL SE

Date of deferred publication of the search report:
 06.05.92 Bulletin 92/19

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Process for the stereochemical inversion of (2S,3S)-2-amino-3-phenyl-1,3-propanediols into their (2R,3R) enantiomers.

(a) A four step process for transforming (2S,3S)-2-amino-3-phenyl-1,3-propanediols into their (2R,3R)-enantiomers is described. The final compounds are useful intermediates for the synthesis of antibiotics like Chloramphenicol, Thiamphenicol and Florfenicol. The starting products generally are discard products in the synthesis of said antibiotics.

EP 90 11 9803

ategory	Citation of document with indica of relevant passage		Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int. CL.5)
D <b>,A</b>	TETRAHEDRON LETTERS, vol. 29, no. 43, 1988, OXF pages 5561 - 5564; C. GIORDANO ET AL.: 'New S Racemization of 2-Amino-1, Intermediates for the Synt Drugs' * the whole document *	ORD GB trategy for 3-Propanediols, Key	1-20	C07B55/00 C07C323/32 C07C317/32 C07C323/39 C07C215/34 C07C213/10 C07D263/06
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	The present search report has been	drawn up for all claims		
	Place of search	Date of completion of the search	750	Emine
X:pa Y:pa	THE HAGUE  CATEGORY OF CITED DOCUMENTS ricularly relevant if taken alone ricularly relevant if combined with another	E : exriter patent d after the filing	ple underlying th ocument, but put date in the applicatio	Eished on, or